



SmartPA Criteria

Proposal

Drug/Drug Class:	Psoriasis Agents, Topical PDL Edit		
First Implementation Date:	May 7, 2008		
Revised Date:	December 8, 2022		
Prepared For:	MO HealthNet		
Prepared By:	MO HealthNet/Conduent		
Criteria Status:	□ Existing Criteria⊠ Revision of Existing Criteria		
	☐ New Criteria		

Executive Summary

Purpose: The MO HealthNet Pharmacy Program will implement a state-specific preferred drug list.

Why Issue Selected:

Psoriasis is a chronic, inflammatory, non-contagious, genetic, immune-mediated, dermatologic condition. The most common type (80-90% of participants) is plaque psoriasis (psoriasis vulgaris) where patches or lesions of skin become inflamed and are covered by a silvery white scale. These plaques frequently occur on the skin of the elbows and knees but can affect any area including the scalp. Psoriasis affects approximately 7.5 million Americans; presentation can occur at any age, but typically occurs between the ages of 15 to 25 years. Psoriasis can range from mild to severe disease and can lead to low self-esteem and depression. During the disease process there is hyperproliferation and abnormal differentiation of the psoriatic epidermis. This disease can also affect the joints and connective tissue, resulting in psoriatic arthritis. Traditionally, pharmacotherapy choices to treat plaque and scalp psoriasis include emollients, topical corticosteroids, vitamin D analogs, calcipotriene/betamethasone, tazarotene, tacrolimus, pimecrolimus, phototherapy, and systemic medications.

Total program savings for the PDL classes will be regularly reviewed.

Program-Specific Information:

;	Preferred Agents	Non-Preferred Agents
•	Calcipotriene Crm/Oint/Soln	 Calcipotriene Foam Calcipotriene/Betamethasone Calcitriol Duobrii[®] Enstilar[®] Sorilux[®] Taclonex[®]

Type of Criteria:	☐ Increased risk of ADE	☑ Preferred Drug List
	☐ Appropriate Indications	☐ Clinical Edit
Data Sources:	☐ Only Administrative Databases	□ Databases + Prescriber-Supplied

Setting & Population

- Drug class for review: Psoriasis Agents, Topical
- Age range: All appropriate MO HealthNet participants

Approval Criteria

- Failure to achieve desired therapeutic outcomes with trial on 1 or more preferred agents in the past 6 months;
 - Documented trial period of preferred agents OR
 - Documented ADE/ADR to preferred agents

Denial Criteria

- Lack of adequate trial on required preferred agents
- Therapy will be denied if all approval criteria are not met

Claim exceeds maximum dosing limitation for the following:

Drug Description	Generic Equivalent	Max Dosing Limitation
DUOBRII 0.01%-0.045% LOT	HALOBETASOL/TAZAROTENE	2 tubes every 28 days

Required Documentation						
Laboratory Results: MedWatch Form:		Progress Notes: Other:				
Disposition of Edit						
Denial: Exception Code "0160" (Preferred Drug List) Rule Type: PDL						
Default Approval Period						

1 year

References

- Evidence-Based Medicine and Fiscal Analysis: "Psoriasis Agents, Topical Therapeutic Class Review", Conduent Business Services, L.L.C., Richmond, VA; January 2022.
- Evidence-Based Medicine Analysis: "Topical Psoriasis Agents", UMKC-DIC; October 2021.
- USPDI, Micromedex; 2022.
- Facts and Comparisons eAnswers (online); 2022 Clinical Drug Information, LLC.

United States and/or other countries.